



Food and Drug Administration
10903 New Hampshire Avenue
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Silver Spring, MD 20993-0002

Resonance Innovations LLC
% Mr. Brian Carter
Quality Assurance Manager
9840 S. 140th Street, Suite 8
OMAHA NE 68138

December 2, 2014

Re: K142755
Trade/Device Name: Siemens 3.0T Blanket Array Coil
Regulation Number: 21 CFR 892.1000
Regulation Name: Magnetic resonance diagnostic device
Regulatory Class: II
Product Code: MOS
Dated: September 24, 2014
Received: September 25, 2014

Dear Mr. Carter:

This letter corrects our substantially equivalent letter of October 23, 2014. We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Industry and Consumer Education at its toll-free number (800) 638 2041 or (301) 796-7100 or at its Internet address

<http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>. Also, please note the regulation entitled, “Misbranding by reference to premarket notification” (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to

<http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH’s Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address

<http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely yours,



for

Janine M. Morris
Director
Division of Radiological Health
Office of In Vitro Diagnostics
and Radiological Health
Center for Devices and Radiological Health

Enclosure

Indications for Use

510(k) Number (if known)

k142755

Device Name

Siemens 3.0T Blanket Array Coil

Indications for Use (Describe)

The coil is indicated for use by the order of a physician to be used as an accessory to a Siemens 3.0T magnetic resonance scanner for general human anatomy imaging as supported by the scanner. These images, when interpreted by a trained physician, may assist in medical diagnosis.

Type of Use (Select one or both, as applicable)

☒ Prescription Use (Part 21 CFR 801 Subpart D)

☐ Over-The-Counter Use (21 CFR 801 Subpart C)

PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON A SEPARATE PAGE IF NEEDED.

FOR FDA USE ONLY

Concurrence of Center for Devices and Radiological Health (CDRH) (Signature)

This section applies only to requirements of the Paperwork Reduction Act of 1995.

DO NOT SEND YOUR COMPLETED FORM TO THE PRA STAFF EMAIL ADDRESS BELOW.

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“An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB number.”

Section 8. 510(k) Summary

Submitter's Name:	Resonance Innovations LLC
Submitter's Address:	9840 South 140 th St., Suite 8 Omaha, NE 68138
Submitter's Telephone:	402-934-2650
Submitter's Contact:	Randall Jones, President
Date 510(k) Summary prepared:	September 24, 2014
Proprietary Name:	Siemens 3.0T Blanket Array Coil Model 516SI3001
Common or Usual Name:	MRI coil(s)
Classification Name:	Coil, Magnetic Resonance, Specialty
Classification Code:	MOS
Predicate Device:	1.5T and 3.0T GE PROCURE™ Array Coil, K140606

Description of the Device

The Siemens 3.0T Blanket Array Coil embeds an MRI antenna set within a quilt-like enclosure providing high-quality images of general human anatomy. It is flexible enough to wrap snugly around small bodies as well as large adult extremities so it can be used to image those challenging regions such as bilateral brachial plexus, bilateral lower extremities, and even heads. This lightweight SemiFlex™ design facilitates effortless and accurate positioning and ensures the multiple antenna elements as close as possible to the target anatomies regardless of patient size. The Blanket Array Coil is an MRI coil contained within a water-resistant case, featuring a soft and cleanable cover. What the patient feels is a lightweight, warm and soothing blanket. What the radiologist sees is a flawless image without compromising comfort. The underlying technology and materials used for manufacture are identical to standard MRI coil technology that has existed for several years. The coil uses similar blocking networks and impedance matching circuits, and it does not transmit energy into the patient, neither predicate nor current submission.

Device Model Number	Device Description
516SI3001	Siemens 3.0T Blanket Array Coil

Indications for Use

The coil is indicated for use by the order of a physician to be used as an accessory to a Siemens 3.0T magnetic resonance scanner for general human anatomy imaging as supported by the scanner. These images, when interpreted by a trained physician, may assist in medical diagnosis.

Technological Characteristics

The comparison between the predicate and the current submission is described, below.

1. **Design.** This submission is for a dedicated coil that may give diagnostic quality images of general human anatomy. It is a multi-channel coil with all channels designed to work at once, receive-only, in conjunction with the system body coil. The predicate multi-channel GE PROCURE™ Array Coil covers similar anatomy, also working as receive-only to the body coil. Both predicate and modified coils have resonant coil elements enclosed in semi-flexible assemblies. The modified device has been slightly modified to integrate with the Siemens MRI scanner. All design principles employed are mature and well-known throughout the industry.
2. **Principles of operation:** The scientific principles of operation (magnetic resonance) are identical between the predicate and modified devices. Theory of operation is very well understood throughout the industry.
3. **Materials.** The same materials are used in the construction of the predicate and modified device. All internal circuitry is encapsulated in flame retardant EVA foam, then completely covered with a nylon fabric or compressed EVA. This nylon fabric has been successfully used for over 10 years in the predicate device with no reported biocompatibility issues. Both modified and predicate devices have been tested for mechanical and electrical safety using IEC60601-1 3rd Edition, which for our application is identical to ES60601-1.
4. **Chemical Composition.** Both predicate and modified devices have a successful biocompatibility track record, as demonstrated by cytotoxicity testing and by their history of use in previously cleared devices.
5. **Energy Source.** This device is a receive-only coil that does not generate its own power, but rather is controlled by the MRI system as the energy source.

Non-Clinical Tests

The predicate and current submission have been subject to similar risk management studies, as listed below, and determined to be substantially equivalent.

1. Blocking network analysis
2. SNR and uniformity analysis
3. Risk management (including hazard analysis and FMEA)
4. Heat testing
5. Compliance testing to IEC60601-1 3rd Edition. IEC60601-1 was chosen for compliance in other world markets. A gap analysis of ES60601-1:2005/(R)2012 and C1:2009/(R)2012 and, a2:2010/(r)2012 with IEC60601-1 was performed and for our devices, the US deviations described by ES60601-1 do not modify the requirements of IEC60601-1. Consequently, we are compliant with ES60601-1.

Clinical Tests

Analyses in all 3 planes (sagittal, coronal, and axial) were run on the Siemens 3.0T Blanket Array Coil to show that the anatomies of the submitted and predicate coils have substantial equivalence.

Substantial Equivalence Decision

As described in this summary, the modified device is substantially equivalent to the predicate device based on the analysis herein. The modified device raises no new concerns of safety or efficacy.